

As  
can't

wherein X is L-serine residue, L-asparagine residue or (S)-2-aminobutyric acid residue and Y is  $\alpha$ -L-amino acid residue, or a salt thereof.

I.  
A3

7. (Amended) A pharmaceutical composition which comprises the compound claimed in claim 1 or its pro-drug and a pharmaceutically acceptable additive.

II.  
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12. (Amended) A pharmaceutical composition which is a gastric mucosa adhesive pharmaceutical composition comprising (a) a compound as claimed in claim 1, (b) a lipid and/or a polyglycerol fatty acid ester and (c) a viscogenic agent capable of being viscous with water.

III.  
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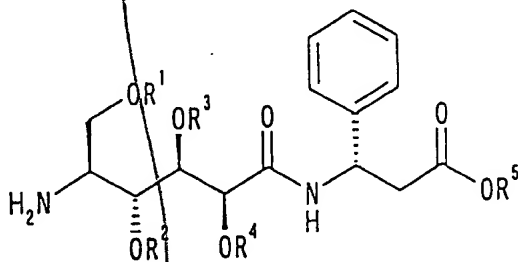
19. (Amended) A method for manufacturing a pharmaceutical composition for *Helicobacter pylori* infectious disease, which comprises mixing the compound according to claim 1 or its pro-drug with a pharmaceutically acceptable additive.

20. (Amended) The method as claimed in claim 19, wherein the composition is for treating or preventing a *Helicobacter pylori* infectious disease.

21. (Amended) The method as claimed in claim 20, wherein the *Helicobacter pylori* infectious disease is gastric or duodenal ulcer, gastritis, gastric cancer or gastric MALT lymphoma.

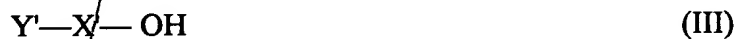
IV.  
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22. (Amended) A method for producing a compound claimed in claim 1, which comprises reacting a compound of the formula:



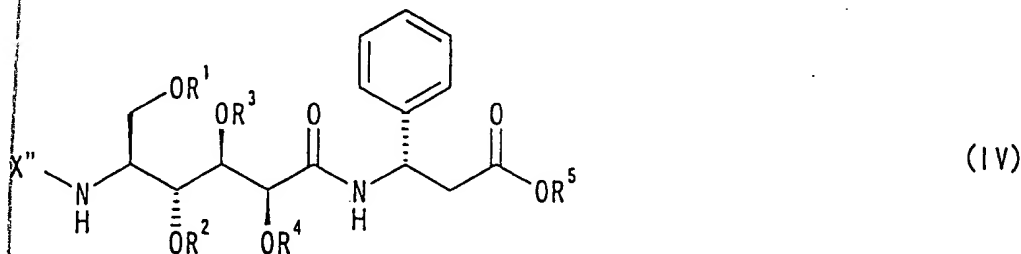
(11)

wherein  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are independently a protecting group for hydroxy group or a hydrogen atom, and  $R^5$  is a protecting group for carboxyl group or a hydrogen atom, a salt thereof or a reactive derivative thereof at the amino group, with a compound of the formula:



wherein  $X'$  is L-serine residue which may be protected, L-asparagine residue which may be protected or (S)-2-aminobutyric acid residue, and  $Y'$  is  $\alpha$ -L-amino acid residue which may be protected, a salt thereof or a reactive derivative thereof at the carboxyl group, if necessary, followed by removing the protecting group.

23. (Amended) A method for producing a compound claimed in claim 1, which comprises reacting a compound of the formula:



wherein  $X''$  is L-serine residue which may be protected, L-asparagine residue which may be protected or (S)-2-aminobutyric acid residue,  $R^1$ ,  $R^2$ ,  $R^3$  and  $R^4$  are independently a protecting group for hydroxy group or a hydrogen atom, and  $R^5$  is a protecting group for carboxyl group or a hydrogen atom, a salt thereof or a reactive derivative thereof at the amino group, with a compound of the formula:



wherein  $Y'$  is  $\alpha$ -L-amino acid residue which may be protected, a salt thereof or a reactive derivative thereof at the carboxyl group, if necessary, followed by removing the protecting group.